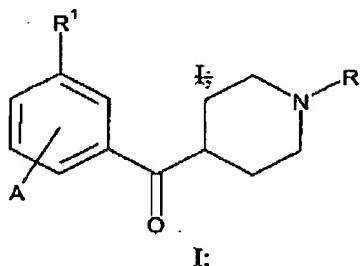


Serial No. 09/890,741

Amendments to the Claims

Claim 1. (Currently amended) A compound of Formula I:



or a pharmaceutical acid addition salt thereof, where;

A is hydrogen, halo, NH₂, or -CF₃;R is hydrogen, C₁-C₄ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, or (C₁-C₆ alkyl)-Ar¹;R¹ is -NH-R²-R³ or -OSO₂-Ar²;Ar, Ar¹, Ar², Ar³, and Ar⁴ are an optionally substituted phenyl or optionally substituted heteroaryl;R² is -CO-, -CS-, or -SO₂-;R³ is ~~hydrogen, optionally substituted C₁-C₆ alkyl, Ar³, NR⁵R⁶, or OR⁵; provided R³ is not hydrogen if R² is either CS or SO₂-;~~~~R⁴ is hydrogen, optionally substituted C₁-C₆ alkyl, or Ar, and~~~~R⁵ and R⁶ are independently hydrogen, optionally substituted C₁-C₆ alkyl, or Ar⁴;~~~~or R⁶ and R⁵ combine, together with the nitrogen atom to which they are attached, to form a pyrrolidine, piperidine, piperazine, 4 substituted piperazine, morpholine or thiomorpholine ring;~~wherein substituted phenyl is phenyl mono-substituted with a substituent selected from the group consisting of halo, nitro, cyano, amino, trifluoromethyl, trifluoromethoxy, phenyl, benzoyl, C₁-C₆ alkyl, C₁-C₆ alkoxy, (C₁-C₄ alkyl)S(O)_n, (C₁-C₄ alkyl)₂ amino, C₁-C₄ acyl, or two or three substituents independently selected from the group consisting of halo, nitro, trifluoromethyl, C₁-C₄ alkyl, and C₁-C₄ alkoxy;

n is 0, 1, or 2;

heteroaryl is an aromatic or benzofused aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur;

Serial No. 09/890,741

substituted heteroaryl is heteroaryl substituted with up to three substituents selected from the group consisting of halo, cyano, nitro, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-;

substituted alkyl is alkyl substituted from 1 to 3 times independently with a substituent selected from the group consisting of halo, hydroxy, phenyl, 2-phenylethylen-1-yl, diphenylmethyl, naphthyl, substituted phenyl, aryloxy, heterocycle, heteroaryloxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₈ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ alkoxy carbonyl, phenyl(C₁-C₄ alkyl), substituted phenyl(C₁-C₄ alkyl), and benzofused C₄-C₈ cycloalkyl; and

heterocycle is aromatic or non-aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur, said ring being optionally benzofused and said ring or benzofused ring being optionally substituted with up to three substituents selected from the groups consisting of halo, C₁-C₄ alkoxy, C₁-C₄ alkyl, cyano, nitro, hydroxy, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-.

Claim 2. (Original) The compound of Claim 1 wherein A is hydrogen.

Claim 3. (Previously presented) The compound of Claim 1 wherein R is methyl.

Claim 4. (Cancelled)

Claim 5. (Cancelled)

Claim 6. (Cancelled)

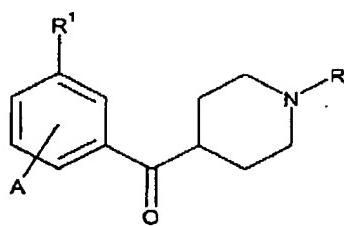
Claim 7. (Currently amended) The compound of Claim 6.1 wherein Ar³ is 4-fluorophenyl.

Claim 8. (Previously presented) The compound of Claim 7 wherein Ar³ is 4-fluorophenyl additionally mono- or disubstituted.

Claim 9. (Previously presented) The compound of Claim 8 wherein Ar³ is selected from the group consisting of 2-iodo-4-fluorophenyl, 2-bromo-4-fluorophenyl, 2-chloro-4-fluorophenyl, 2,4-difluorophenyl, and 2-methyl-4-fluorophenyl, and 2,4,6-trifluorophenyl.

Serial No. 09/890,741

Claim 10. (Currently amended) A pharmaceutical formulation comprising a compound of I:



I;

or a pharmaceutical acid addition salt thereof, where;

A is hydrogen, halo, NH₂, or -CF₃;

R is hydrogen, C₁-C₄ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, or (C₁-C₆ alkyl)-Ar¹;

R¹ is -NH-R²-R³ or ~~OSO₂Ar²~~;

Ar, Ar¹, Ar², Ar³, and Ar⁴ are an optionally substituted phenyl or optionally substituted heteroaryl;

R² is -CO-, -CS-, or ~~SO₂-~~;

R³ is ~~hydrogen, optionally substituted C₁-C₆ alkyl, Ar³, NR⁵R⁶, or OR⁵; provided R³ is not hydrogen if R² is either CS- or SO₂-~~;

~~R⁴ is hydrogen, optionally substituted C₁-C₆ alkyl, or Ar; and R⁵ and R⁶ are independently hydrogen, optionally substituted C₁-C₆ alkyl, or Ar⁴; or R⁶ and R⁵ combine, together with the nitrogen atom to which they are attached, to form a pyrrolidine, piperidine, piperezine, 4 substituted piperazine, morpholine or thiomorpholine ring;~~

wherein substituted phenyl is phenyl mono-substituted with a substituent selected from the group consisting of halo, nitro, cyano, amino, trifluoromethyl, trifluoromethoxy, phenyl, benzoyl, C₁-C₆ alkyl, C₁-C₆ alkoxy, (C₁-C₄ alkyl)S(O)_n, (C₁-C₄ alkyl)₂ amino, C₁-C₄ acyl, or two or three substituents independently selected from the group consisting of halo, nitro, trifluoromethyl, C₁-C₄ alkyl, and C₁-C₄ alkoxy;

n is 0, 1, or 2;

heteroaryl is an aromatic or benzofused aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur;

Serial No. 09/890,741

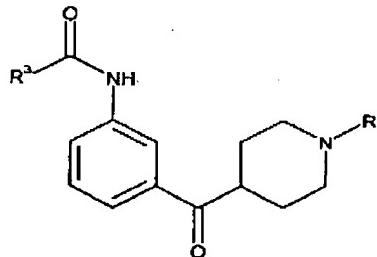
substituted heteroaryl is heteroaryl substituted with up to three substituents selected from the group consisting of halo, cyano, nitro, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n;-

substituted alkyl is alkyl substituted from 1 to 3 times independently with a substituent selected from the group consisting of halo, hydroxy, phenyl, 2-phenylethylen-1-yl, diphenylmethyl, naphthyl, substituted phenyl, aryloxy, heterocycle, heteroaryloxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₈ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ alkoxy carbonyl, phenyl(C₁-C₄ alkyl), substituted phenyl(C₁-C₄ alkyl), and benzofused C₄-C₈ cycloalkyl; and

heterocycle is aromatic or non-aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur, said ring being optionally benzofused and said ring or benzofused ring being optionally substituted with up to three substituents selected from the groups consisting of halo, C₁-C₄ alkoxy, C₁-C₄ alkyl, cyano, nitro, hydroxy, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-.

Claims 11 - 13. (Canceled)

Claim 14. (Previously presented) A process of making the compounds of formula I(a):



I(a)

wherein R³ is hydrogen, optionally substituted C₁-C₆ alkyl, Ar³, -NR⁵R⁶, or OR⁵;

R⁵ and R⁶ are independently hydrogen, optionally substituted C₁-C₈ alkyl, or Ar⁴; or R⁶ and R⁵ combine, together with the nitrogen atom to which they are attached, to form a pyrrolidine, piperidine, piperazine, 4-substituted piperazine, morpholine or thiomorpholine ring; and

Ar³ and Ar⁴ are independently an optionally substituted phenyl or optionally substituted heteroaryl;

wherein substituted phenyl is phenyl mono-substituted with a substituent selected from the group consisting of halo, nitro, cyano, amino, trifluoromethyl, trifluoromethoxy,

Serial No. 09/890,741

phenyl, benzoyl, C₁-C₆ alkyl, C₁-C₆ alkoxy, (C₁-C₄ alkyl)S(O)_n, (C₁-C₄ alkyl)₂ amino, C₁-C₄ acyl, or two or three substituents independently selected from the group consisting of halo, nitro, trifluoromethyl, C₁-C₄ alkyl, and C₁-C₄ alkoxy;

n is 0, 1, or 2;

heteroaryl is an aromatic or benzofused aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur;

substituted heteroaryl is heteroaryl substituted with up to three substituents independently selected from the group consisting of halo, cyano, nitro, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-;

substituted alkyl is alkyl substituted from 1 to 3 times independently with a substituent selected from the group consisting of halo, hydroxy, phenyl, 2-phenylethylen-1-yl, diphenylmethyl, naphthyl, substituted phenyl, aryloxy, heterocycle, heteroaryloxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₈ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ alkoxy carbonyl, phenyl(C₁-C₄ alkyl), substituted phenyl(C₁-C₄ alkyl), and benzofused C₄-C₈ cycloalkyl;

heterocycle is aromatic or non-aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur, said ring being optionally benzofused and said ring or benzofused ring being substituted with up to three substituents selected independently from the groups consisting of halo, C₁-C₄ alkoxy, C₁-C₄ alkyl, cyano, nitro, hydroxy, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-;

comprising:

- (a) protecting 4-benzoylpiperidine hydrochloride to form an N-protected 4-benzoylpiperidine hydrochloride;
- (b) nitrating the N-protected 4-benzoylpiperidine hydrochloride to form a mixture of N-protected 4-(mono nitrobenzoyl)piperidines;
- (c) deprotecting the N-protected 4-(mononitrobenzoyl)-piperidine mixture to form a mixture of 4-(mononitrobenzoyl)piperidines;
- (d) separating the 4-(3-nitrobenzoyl)piperidine from the mixture of 4-(mononitrobenzoyl)piperidines;
- (e) reducing the 4-(3-nitrobenzoyl)piperidine to form 4-(3-aminobenzoyl)piperidine; and
- (f) acylating the 4-(3-aminobenzoyl)piperidine.

Claim 15. (Original) The process of Claim 14 wherein steps a) and b) are combined.

Serial No. 09/890,741

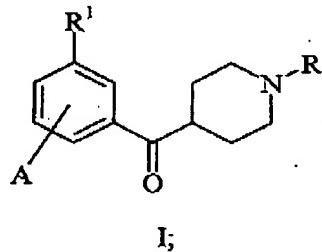
Claim 16. (Previously presented) The process of Claim 14 wherein the source of the protecting group of step a) is trifluoroacetic anhydride.

Claim 17. (Previously presented) The process of Claim 14 wherein the N-protected 4-benzoylpiperidine hydrochloride is nitrated with ammonium nitrate.

Claim 18. (Previously presented) The process of Claim 16 wherein the N-protected 4-benzoylpiperidine hydrochloride is nitrated with ammonium nitrate.

Claim 19. (Canceled)

Claim 20. (Previously presented) A method for treating migraine in a mammal comprising administering to a mammal in need of such treatment an effective amount of a compound of formula I:



or a pharmaceutical acid addition salt thereof, where;

A is hydrogen, halo, NH₂, or -CF₃;

R is hydrogen, C₁-C₄ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, or (C₁-C₆ alkyl)-Ar¹;

R¹ is -NH-R²-R³ or -OSO₂Ar²;

Ar, Ar¹, Ar², Ar³, and Ar⁴ are an optionally substituted phenyl or optionally substituted heteroaryl;

R² is -CO-, -CS-, or -SO₂-;

R³ is hydrogen, optionally substituted C₁-C₆ alkyl, Ar³, -NR⁵R⁶, or OR⁵; provided R³ is not hydrogen if R² is either -CS- or -SO₂-;

R⁴ is hydrogen, optionally substituted C₁-C₆ alkyl, or Ar, and

Serial No. 09/890,741

R⁵ and R⁶ are independently hydrogen, optionally substituted C₁-C₈ alkyl, or Ar⁴; or R⁶ and R⁵ combine, together with the nitrogen atom to which they are attached, to form a pyrrolidine, piperidine, piperazine, 4-substituted piperazine, morpholine or thiomorpholine ring;

wherein substituted phenyl is phenyl mono-substituted with a substituent selected from the group consisting of halo, nitro, cyano, amino, trifluoromethyl, trifluoromethoxy, phenyl, benzoyl, C₁-C₆ alkyl, C₁-C₆ alkoxy, (C₁-C₄ alkyl)S(O)_n, (C₁-C₄ alkyl)₂ amino, C₁-C₄ acyl, or two or three substituents independently selected from the group consisting of halo, nitro, trifluoromethyl, C₁-C₄ alkyl, and C₁-C₄ alkoxy;

n is 0, 1, or 2;

heteroaryl is an aromatic or benzofused aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur;

substituted heteroaryl is heteroaryl substituted with up to three substituents selected from the group consisting of halo, cyano, nitro, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-,

substituted alkyl is alkyl substituted from 1 to 3 times independently with a substituent selected from the group consisting of halo, hydroxy, phenyl, 2-phenylethylen-1-yl, diphenylmethyl, naphthyl, substituted phenyl, aryloxy, heterocycle, heteroaryloxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₈ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ alkoxy carbonyl, phenyl(C₁-C₄ alkyl), substituted phenyl(C₁-C₄ alkyl), and benzofused C₄-C₈ cycloalkyl; and

heterocycle is aromatic or non-aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur, said ring being optionally benzofused and said ring or benzofused ring being optionally substituted with up to three substituents selected from the groups consisting of halo, C₁-C₄ alkoxy, C₁-C₄ alkyl, cyano, nitro, hydroxy, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-.

Claim 21. (Previously presented) The method according to Claim 20 where the mammal is a human.

Claim 22. (Canceled)

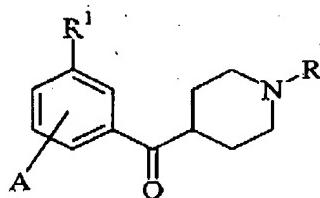
Claim 23. (Currently amended) The compound of Claim 61 where A is hydrogen and R is methyl.

Serial No. 09/890,741

Claim 24. (Previously presented) The compound of Claim 7 where A is hydrogen and R is methyl.

Claim 25. (Currently amended) The compound of Claim 6 1 where R¹ is -NH-R²-R³, R² is C=O and R³ is substituted halophenyl.

Claim 26. (Currently amended) A method for activating 5-HT_{1F} receptors in mammals comprising administering to a mammal in need of such activation an effective amount of a compound of formula I:



I;

or a pharmaceutical acid addition salt thereof, where;

A is hydrogen, halo, NH₂, or -CF₃;

R is hydrogen, C₁-C₄ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl, or (C₁-C₆ alkyl)-Ar¹;

R¹ is -NH-R²-R³ or ~~OSO₂Ar²~~;

Ar, Ar¹, Ar², Ar³, and Ar⁴ are an optionally substituted phenyl or optionally substituted heteroaryl;

R² is -CO-, -CS-, or ~~SO₂-~~;

R³ is hydrogen, optionally substituted C₁-C₆ alkyl, Ar³, NR⁵R⁶, or OR⁵, provided R³ is not hydrogen if R² is either CS or SO₂;

R⁴ is hydrogen, optionally substituted C₁-C₆ alkyl, or Ar; and

R⁵ and R⁶ are independently hydrogen, optionally substituted C₁-C₆ alkyl, or Ar⁴; or R⁶ and R⁵ combine together with the nitrogen atom to which they are attached, to form a pyrrolidine, piperidine, piperazine, 4-substituted piperazine, morpholine or thiomorpholine ring;

wherein substituted phenyl is phenyl mono-substituted with a substituent selected from the group consisting of halo, nitro, cyano, amino, trifluoromethyl, trifluoromethoxy, phenyl, benzoyl, C₁-C₆ alkyl, C₁-C₆ alkoxy, (C₁-C₄ alkyl)S(O)_n, (C₁-C₄ alkyl)₂ amino, C₁-C₄

Serial No. 09/890,741

acyl, or two or three substituents independently selected from the group consisting of halo, nitro, trifluoromethyl, C₁-C₄ alkyl, and C₁-C₄ alkoxy;

n is 0, 1, or 2;

heteroaryl is an aromatic or benzofused aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur;

substituted heteroaryl is heteroaryl substituted with up to three substituents selected from the group consisting of halo, cyano, nitro, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-;

substituted alkyl is alkyl substituted from 1 to 3 times independently with a substituent selected from the group consisting of halo, hydroxy, phenyl, 2-phenylethylen-1-yl, diphenylmethyl, naphthyl, substituted phenyl, aryloxy, heterocycle, heteroaryloxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₈ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ alkoxycarbonyl, phenyl(C₁-C₄ alkyl), substituted phenyl(C₁-C₄ alkyl), and benzofused C₄-C₈ cycloalkyl; and

heterocycle is aromatic or non-aromatic 5 or 6 membered ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur, said ring being optionally benzofused and said ring or benzofused ring being optionally substituted with up to three substituents selected from the groups consisting of halo, C₁-C₄ alkoxy, C₁-C₄ alkyl, cyano, nitro, hydroxy, (C₁-C₄ alkyl)-S(O)_n-, and phenyl-S(O)_n-.

Claim 27. (Previously presented) The method according to Claim 26 where the mammal is a human.

Claims 28. (Cancelled)

Claim 29. (Cancelled)